4. (Twice Amended) The pharmaceutical composition of Claim 1, wherein said hydroxyethylstarch contains at least 10% hydroxyethylstarch with a molecular weight of 25,000--45,000 atomic mass units.

5. (Amended) The pharmaceutical composition of Claim 1, wherein said gelatin derivatives have a molecular weight of 20,000~35,000 atomic mass units, and are selected from urea--conjugated gelatin, modified liquid gelatin, oxidized polygelatin and degraded gelatin polypeptide.

6. (Amended) The pharmaceutical composition of Claim 1, wherein said dextran has a molecular weight of 40,000--230,000 atomic mass units; said carboxymethylstarch has a molecular weight of 30,000--80,000 atomic mass units, said PVP has a molecular weight of 5,000--700,000 atomic mass units, said condensed glucose has a molecular weight of 8,000~12,000 atomic mass units, said sodium alginate has a molecular weight of 20,000--26,000 atomic mass units, said pectin has a molecular weight of 20,000~40,000 atomic mass units; and said pentahydroxyethylstarch has a molecular weight of about 264,000 atomic mass units.

7. (Amended) A method for preparing the pharmaceutical composition of Claim 1, comprising:

dissolving 3--18g of one or more substances selected from hydroxyethylstarch, dextran, carboxymethylstarch, PVP, gelatin derivatives, condensed glucose, glucose. fructose, lactose, glycerin, xylitol, sodium alginate, N--2--hydroxypropylacrylamide, ethylene epoxide-polypropylene glycol, pectin, and pentahydroxyethylstarch, in total of 100 ml of one injection or mixture of several injections selected from water, physiological saline, balanced buffers, glucose solution, sodium lactate solution, sodium acetate solution, Tris solution, and glucose and sodium chloride solution;

adding 1.5g sodium chloride and 0—5.4g of one or more substances selected from sodium chloride. sodium bicarbonate, potassium chloride, magnesium sulfate, calcium chloride, calcium gluconate, calcium lactate, sodium lactate, sodium acetate, and Tris,

mixing to dissolution to obtain said pharmaceutical composition.

Remarks

The applicant voluntarily made the foregoing amendments to increase the clarity of the claims. It is respectfully submitted that the amendments are supported by the specification as filed, so that no new matter has been added. A notice of allowance of the pending claims is respectfully solicited.

Respectfully submitted,

3/18/02
Date

By _____

Alvin R. Wirthlin, Reg. No. 40, 267
Akin, Gump, Strauss, Hauer & Feld, L.L.P.
P.O. Box 688
Dallas, TX 75313-0688
(214) 969-2800